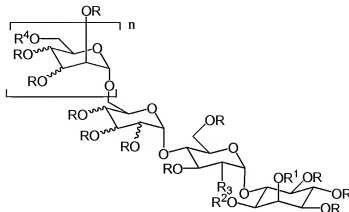


*In the Claims:*

1. **(previously presented)** A compound represented by formula I:



I

wherein,

n is 1, 3, or 4;

R represents independently for each occurrence H, alkyl, aryl, -CH<sub>2</sub>-aryl, -C(O)-alkyl, -C(O)-aryl, or -Si(alkyl)<sub>3</sub>;

R<sup>1</sup> and R<sup>2</sup> are independently H, -CH<sub>2</sub>-aryl, -C(O)-alkyl, -C(O)-aryl, -Si(alkyl)<sub>3</sub>; or R<sup>1</sup> and R<sup>2</sup> taken together are C(CH<sub>3</sub>)<sub>2</sub>, P(O)OH, or P(O)OR<sup>5</sup>;

R<sup>3</sup> is amino, -N<sub>3</sub>, or -NH<sub>3</sub>X;

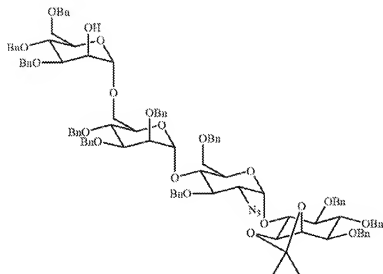
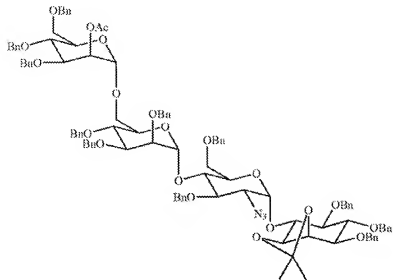
R<sup>4</sup> represents independently for each occurrence alkyl, aryl, -CH<sub>2</sub>-aryl, -C(O)-alkyl, -C(O)-aryl, -Si(alkyl)<sub>3</sub>, or -P(O)(OR<sup>5</sup>)<sub>2</sub>;

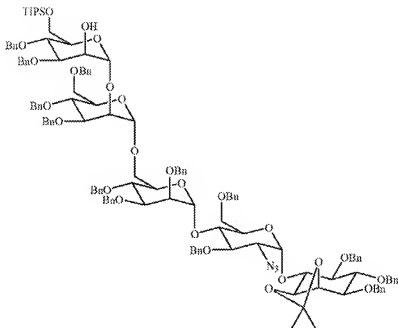
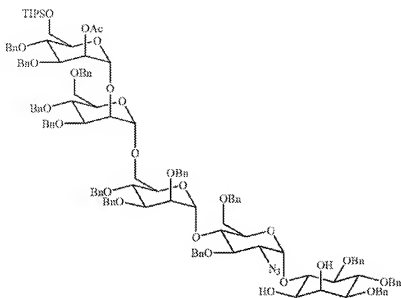
R<sup>5</sup> represents independently for each occurrence H, Li<sup>+</sup>, Na<sup>+</sup>, K<sup>+</sup>, Rb<sup>+</sup>, Cs<sup>+</sup>, aryl, or an optionally substituted alkyl group; and

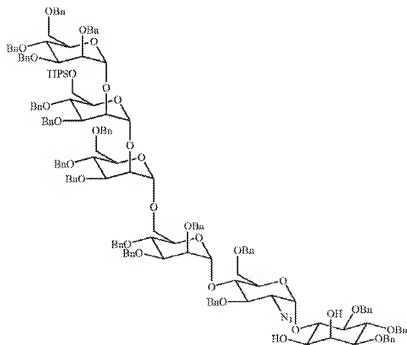
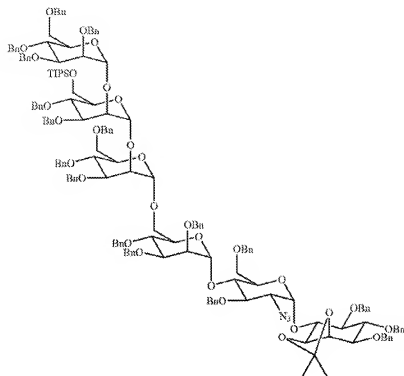
X is a halogen, alkyl carboxylate, or aryl carboxylate.

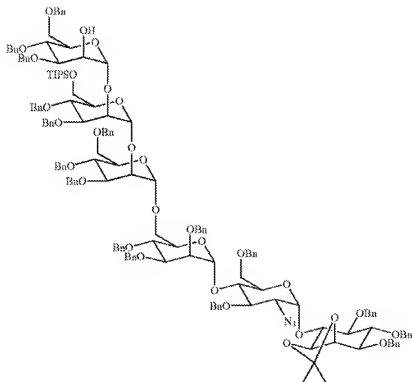
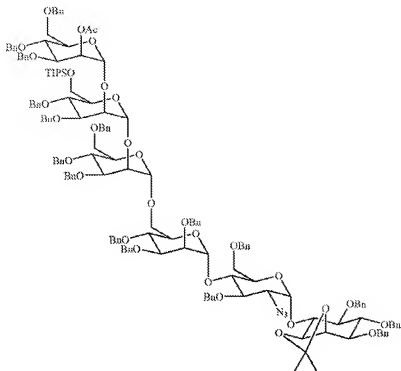
2. **(canceled)**
3. **(original)** The compound of claim 1, wherein n is 3.
4. **(original)** The compound of claim 1, wherein R is H.

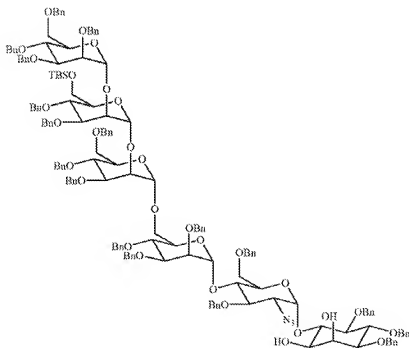
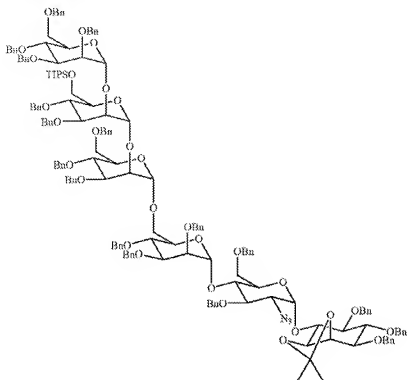
5. **(original)** The compound of claim 1, wherein  $R^1$  and  $R^2$  taken together are  $P(O)OR^5$ .
6. **(original)** The compound of claim 1, wherein  $R^3$  is  $N_3$ .
7. **(original)** The compound of claim 1, wherein  $R^3$  is  $-NH_3X$ .
8. **(previously presented)** The compound of claim 1, wherein  $R^4$  represents independently for each occurrence  $-CH_2Ph$ , or  $-Si(alkyl)_3$ .
9. **(previously presented)** The compound of claim 1, wherein  $R^4$  represents independently for each occurrence  $-CH_2Ph$ , -or  $P(O)OR^5$ ; and  $R^5$  is an optionally substituted alkyl group.
10. **(currently amended)** A compound selected from the group consisting of:

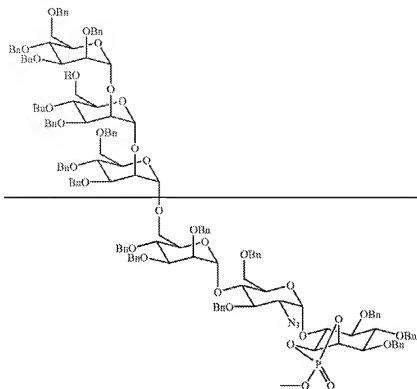




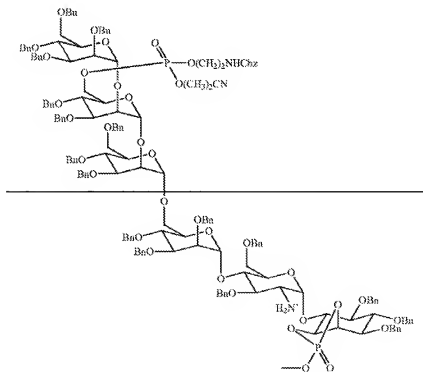
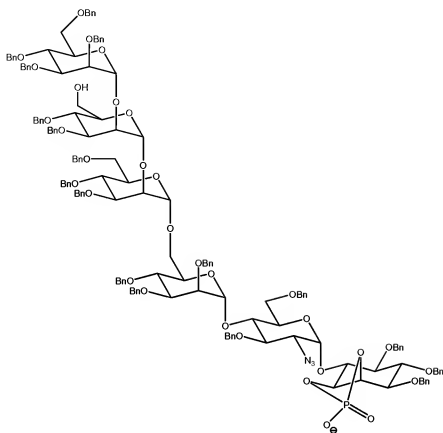


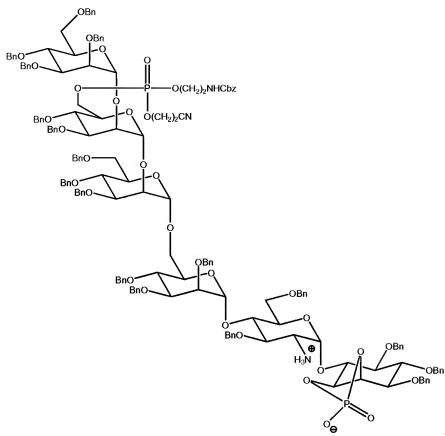


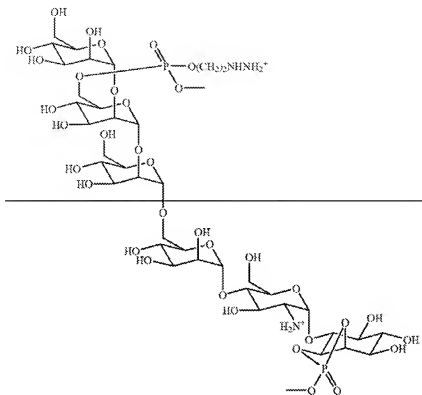


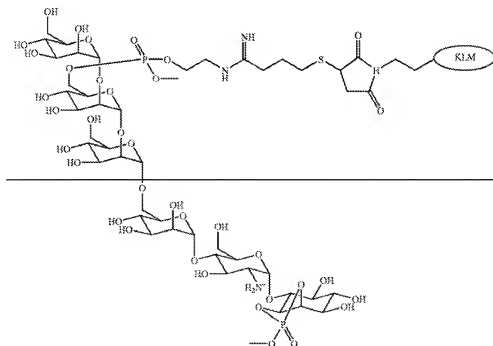
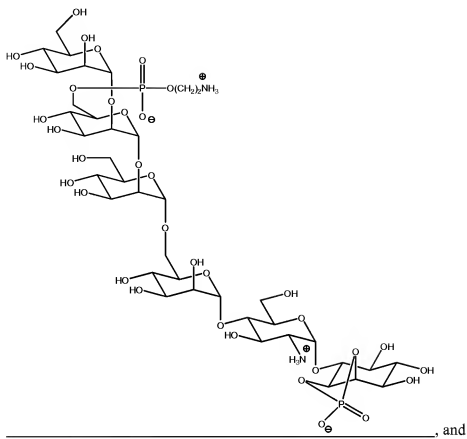


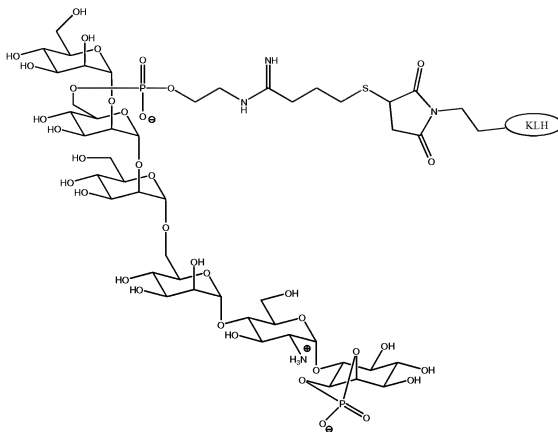




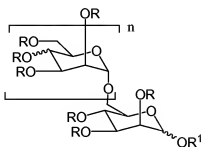








11. (previously presented) A compound represented by formula II:



II

wherein,

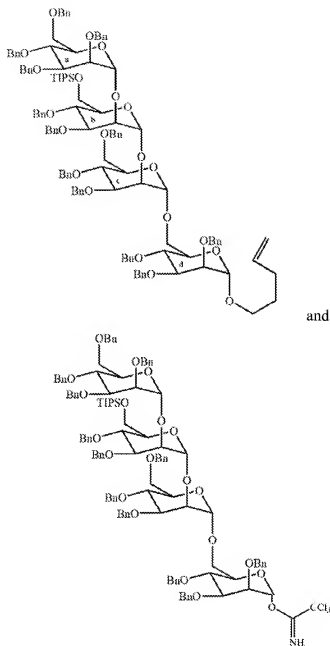
n is 3, or 4;

R represents independently for each occurrence H, alkyl, aryl,  $-\text{CH}_2\text{-aryl}$ ,  $-\text{C}(\text{O})\text{-alkyl}$ ,  $-\text{C}(\text{O})\text{-aryl}$ , or  $-\text{Si}(\text{alkyl})_3$ ;

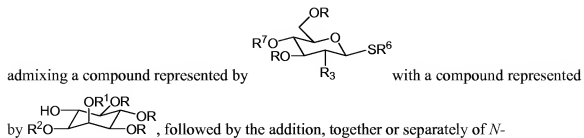
$\text{R}^1$  is  $-(\text{CH}_2)_m\text{CH}=\text{CH}_2$  or trichloroacetimidate; and

m is 1-6.

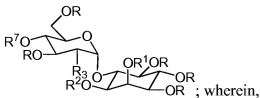
12. **(canceled)**
13. **(original)** The compound of claim 11, wherein n is 3.
14. **(original)** The compound of claim 11, wherein m is 3.
15. **(original)** The compound of claim 11, wherein R represents independently for each occurrence -CH<sub>2</sub>-aryl or -Si(alkyl)<sub>3</sub>.
16. **(original)** The compound of claim 11, wherein R represents independently for each occurrence benzyl or -Si(iPr)<sub>3</sub>.
17. **(previously presented)** The compound of claim 11, wherein R<sup>1</sup> is trichloroacetimidate and R represents independently for each occurrence benzyl or -Si(iPr)<sub>3</sub>.
18. **(previously presented)** The compound of claim 11, wherein said compound of formula **II** is selected from the group consisting of:



19. (previously presented) A method comprising the step of:



iodosuccinimide and silver triflate, thereby forming a compound represented by



R represents independently for each occurrence H, alkyl, aryl,  $-\text{CH}_2\text{-aryl}$ ,  $-\text{C}(\text{O})\text{-alkyl}$ ,  $-\text{C}(\text{O})\text{-aryl}$ , or  $-\text{Si}(\text{alkyl})_3$ ;

$\text{R}^1$  and  $\text{R}^2$  are independently H,  $-\text{CH}_2\text{-aryl}$ ,  $-\text{C}(\text{O})\text{-alkyl}$ ,  $-\text{C}(\text{O})\text{-aryl}$ ,  $-\text{Si}(\text{alkyl})_3$ ; or  $\text{R}^1$  and  $\text{R}^2$  taken together are  $\text{C}(\text{CH}_3)_2$ ,  $\text{P}(\text{O})\text{OH}$ , or  $\text{P}(\text{O})\text{OR}^5$ ;

$\text{R}^3$  is amino,  $-\text{N}_3$ , or  $-\text{NH}_3\text{X}$ ;

$\text{R}^5$  represents independently for each occurrence H,  $\text{Li}^+$ ,  $\text{Na}^+$ ,  $\text{K}^+$ ,  $\text{Rb}^+$ ,  $\text{Cs}^+$ , aryl, or an optionally substituted alkyl group;

$\text{R}^6$  is alkyl or aryl;

$\text{R}^7$  is alkyl, aryl,  $-\text{CH}_2\text{-aryl}$ ,  $-\text{C}(\text{O})\text{-alkyl}$ ,  $-\text{C}(\text{O})\text{-aryl}$ , or  $-\text{Si}(\text{alkyl})_3$ ; and

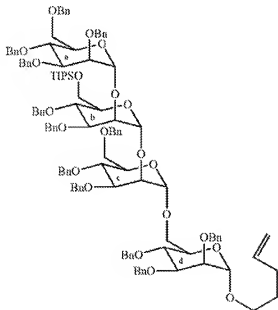
X is a halogen, alkyl carboxylate, or aryl carboxylate.

20. **(original)** The method of claim 19, wherein R is  $-\text{CH}_2\text{-aryl}$ .
21. **(original)** The method of claim 19, wherein  $\text{R}^1$  and  $\text{R}^2$  taken together are  $\text{C}(\text{CH}_3)_2$ .
22. **(original)** The method of claim 19, wherein  $\text{R}^3$  is  $-\text{N}_3$ .
23. **(original)** The method of claim 19, wherein  $\text{R}^6$  is alkyl.
24. **(original)** The method of claim 19, wherein  $\text{R}^7$  is  $-\text{C}(\text{O})\text{-alkyl}$ .
25. **(original)** The method of claim 19, wherein R is benzyl,  $\text{R}^1$  and  $\text{R}^2$  taken together are  $\text{C}(\text{CH}_3)_2$ , and  $\text{R}^3$  is  $-\text{N}_3$ .
26. **(original)** The method of claim 19, wherein R is benzyl,  $\text{R}^1$  and  $\text{R}^2$  taken together are  $\text{C}(\text{CH}_3)_2$ ,  $\text{R}^3$  is  $-\text{N}_3$ , and  $\text{R}^6$  is ethyl.
27. **(previously presented)** A method of preparing a tetrasaccharide, comprising the steps of:

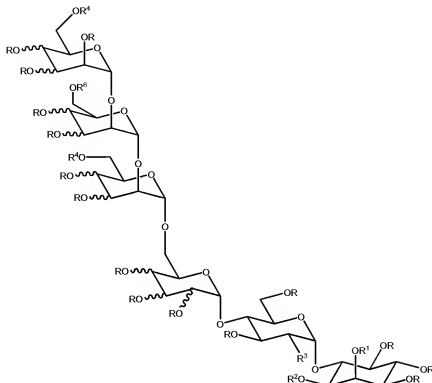


covalently binding a mannopyranoside to a solid support to provide a first substrate, reacting said first substrate with a mannopyranose trichloroacetimidate to give a disaccharide bound to said solid support, reacting said disaccharide with a mannopyranose trichloroacetimidate to give a trisaccharide bound to said solid support, reacting said trisaccharide with a mannopyranose trichloroacetimidate to give a tetrasaccharide bound to said solid support, and cleaving said tetrasaccharide from said solid support.

28. **(original)** The method of claim 27, wherein said mannopyranoside is bound to said solid support through a glycosidic linkage.
29. **(original)** The method of claim 27, wherein said tetrasaccharide is cleaved from said solid support using Grubbs' catalyst.
30. **(previously presented)** The method of claim 27, wherein said tetrasaccharide is



31. (new) A compound represented by formula I:



I

wherein,

R represents independently for each occurrence H, alkyl, aryl,  $-CH_2$ -aryl,  $-C(O)$ -alkyl,  $-C(O)$ -aryl, or  $-Si(alkyl)_3$ ;

$R^1$  and  $R^2$  are independently H,  $-CH_2$ -aryl,  $-C(O)$ -alkyl,  $-C(O)$ -aryl,  $-Si(alkyl)_3$ ; or  $R^1$  and  $R^2$  taken together are  $C(CH_3)_2$ ,  $P(O)OH$ , or  $P(O)OR^5$ ;

$R^3$  is amino,  $-N_3$ , or  $-NH_3X$ ;

$R^4$  represents independently for each occurrence H, alkyl, aryl,  $-CH_2$ -aryl,  $-C(O)$ -alkyl,  $-C(O)$ -aryl,  $-Si(alkyl)_3$ , or  $-P(O)(OR^5)_2$ ;

$R^5$  represents independently for each occurrence H,  $Li^+$ ,  $Na^+$ ,  $K^+$ ,  $Rb^+$ ,  $Cs^+$ , aryl, or an optionally substituted alkyl group; and

$R^6$  represents independently for each occurrence alkyl, aryl,  $-CH_2$ -aryl,  $-C(O)$ -alkyl,  $-C(O)$ -aryl,  $-Si(alkyl)_3$ , or  $-P(O)(OR^5)_2$ ;

X is a halogen, alkyl carboxylate, or aryl carboxylate.

32. (new) The compound of claim 31, wherein R is H.
33. (new) The compound of claim 31, wherein  $R^1$  and  $R^2$  taken together are  $P(O)OR^5$ .
34. (new) The compound of claim 31, wherein  $R^3$  is  $-NH_3X$ .
35. (new) The compound of claim 31, wherein  $R^4$  is H.
36. (new) The compound of claim 31, wherein  $R^6$  is  $-P(O)(OR^5)_2$ .
37. (new) The compound of claim 31, wherein R is H;  $R^1$  and  $R^2$  taken together are  $P(O)OR^5$ ;  $R^3$  is  $-NH_3X$ ;  $R^4$  is H; and  $R^6$  is  $-P(O)(OR^5)_2$ .